

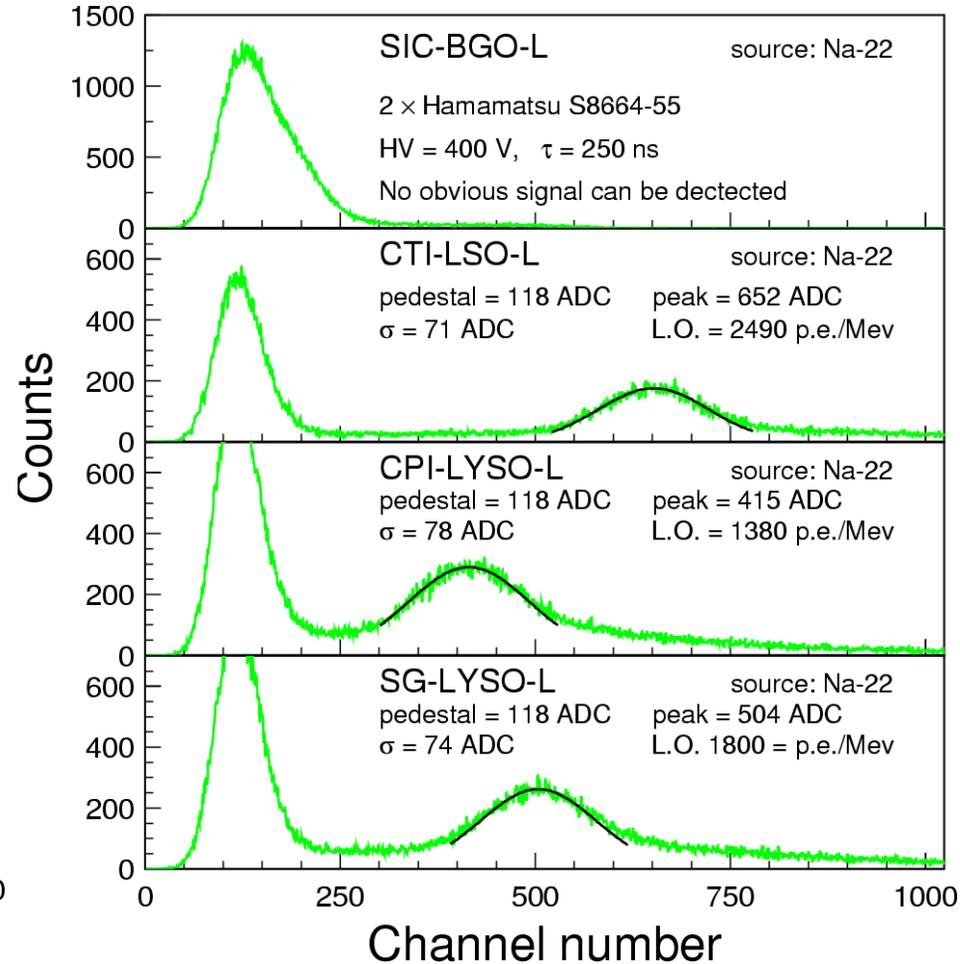
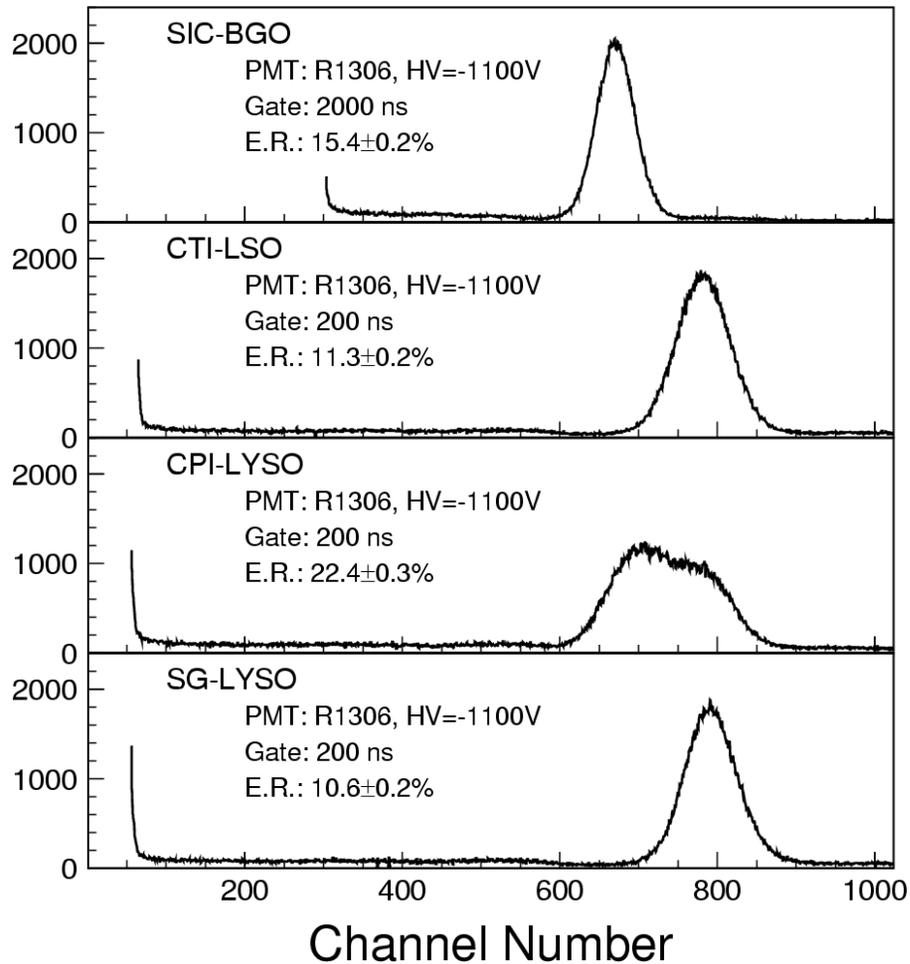


Emission Spectra of LSO and LYSO Crystal Scintillators Excited by UV Light, X-ray and γ -ray

Liyuan Zhang, Rihua Mao and Ren-yuan Zhu

California Institute of Technology

We reported at Puerto Rico (NSS05, N12-6) a comparison of light output of large size (2.5 x 2.5 x 20 cm) LSO and LYSO samples, and found that a CTI LSO has a higher light output with APD readout, but not with PMT readout. This anomaly disappeared after γ -ray irradiation to 1 Mrad.

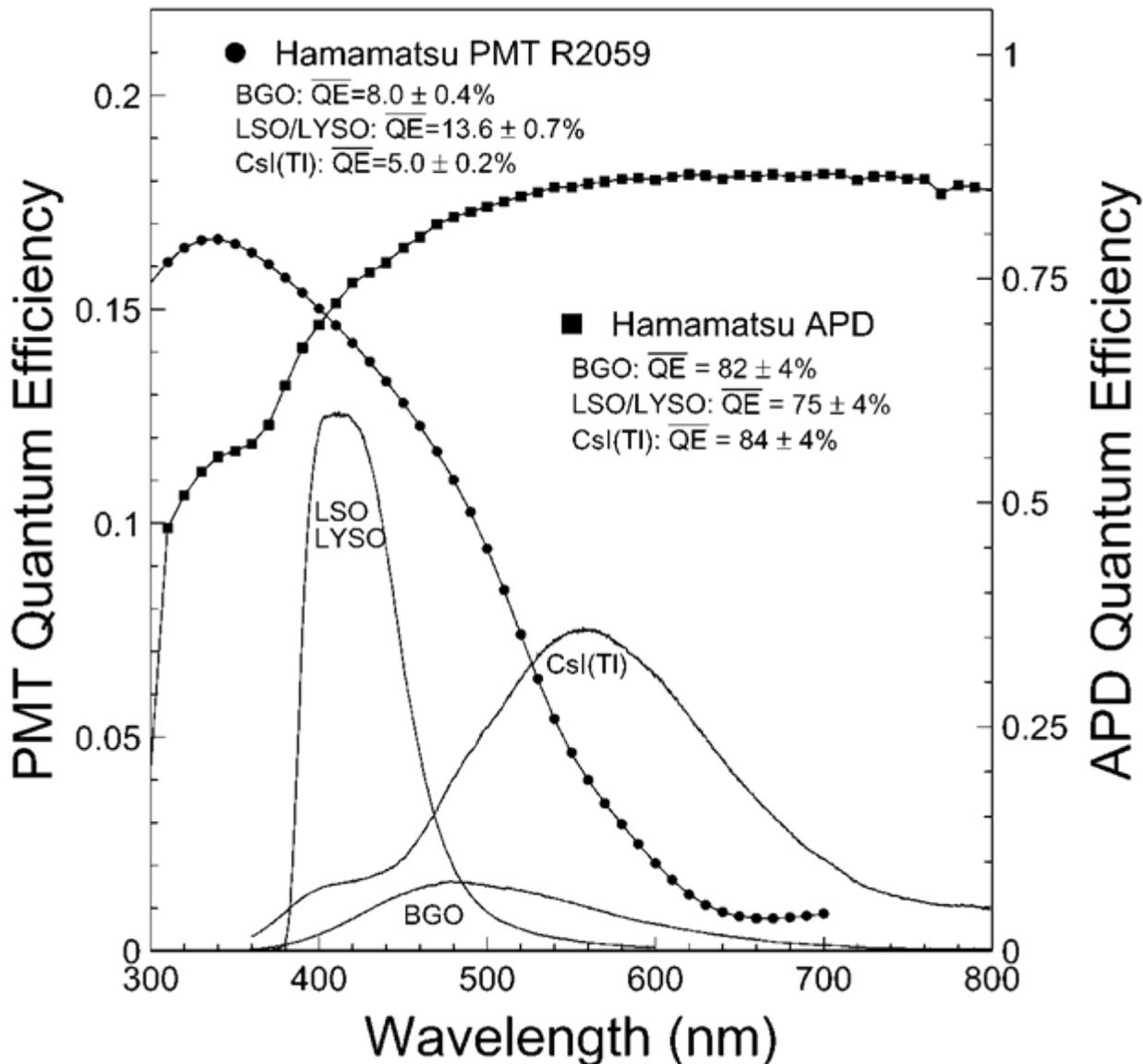


Quantum efficiency and Emission

Why?

The LSO sample had an emission peaked at longer wavelength under γ -ray excitation, and the emission was changed after γ -ray irradiations.

Can we prove it?



Six Large LSO and LYSO Samples

2.5 x 2.5 x 20 cm (18 X₀) Bar

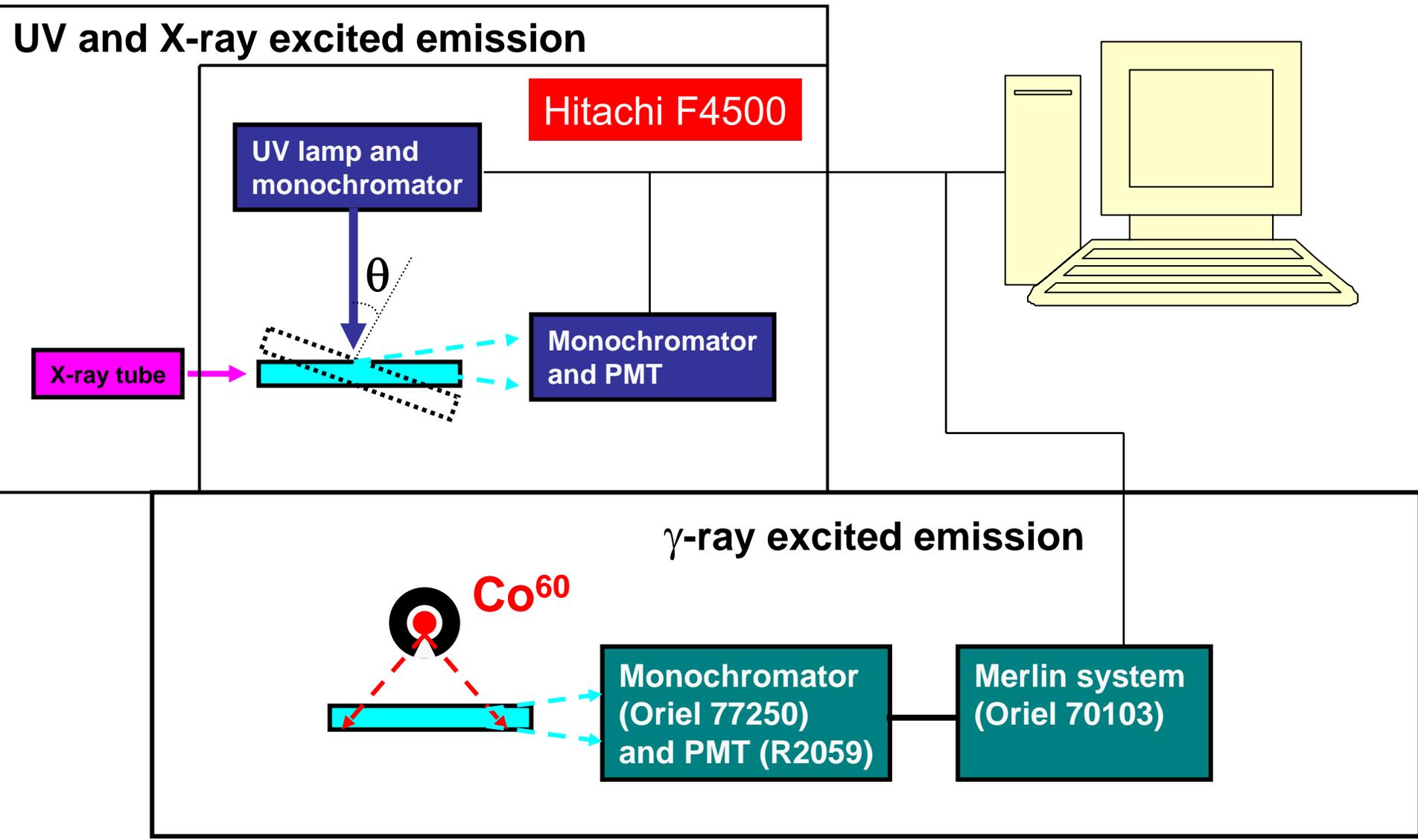


Three CTI LSO samples are provided by Chuck Melcher.

Three LYSO samples are purchased from Saint-Gobain.

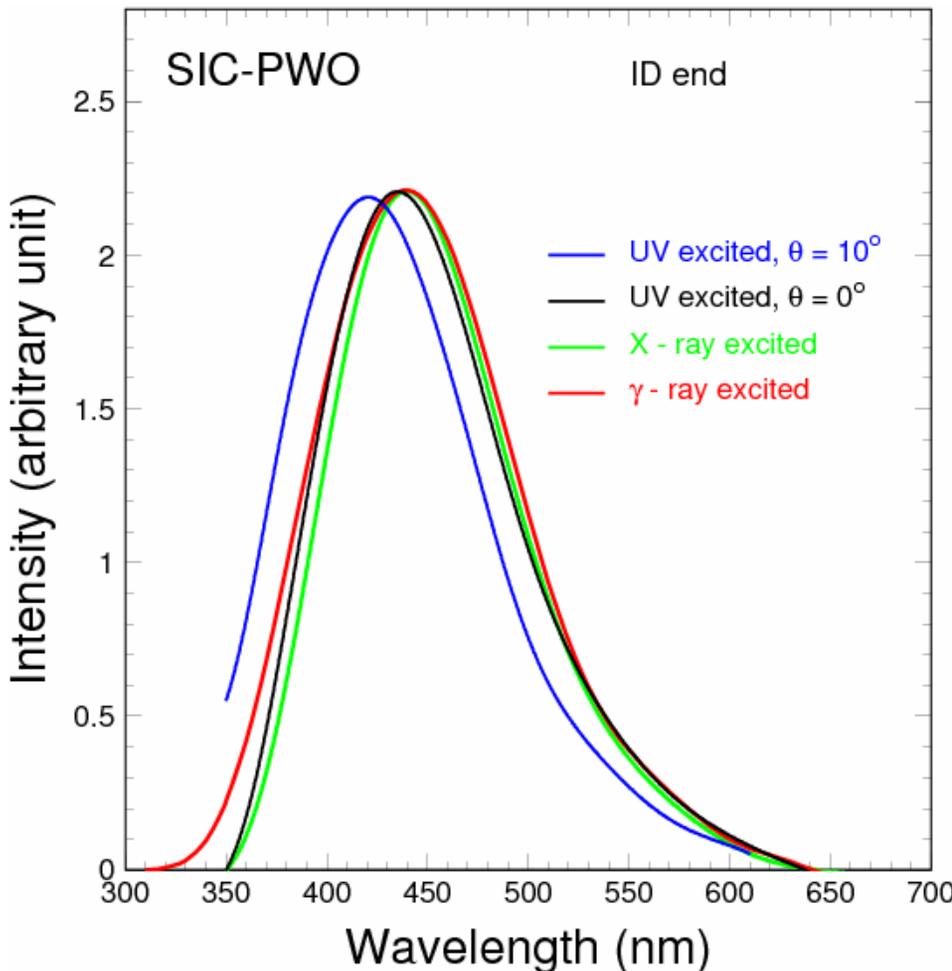
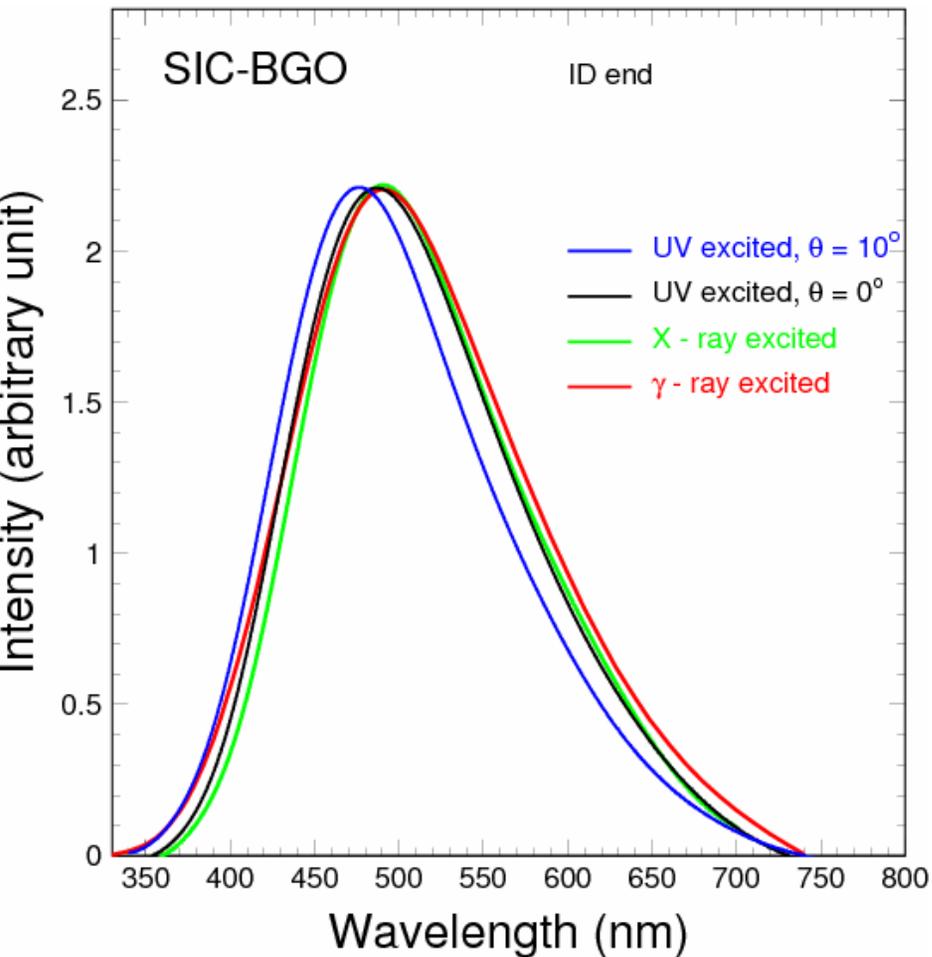
UV, X-ray & γ -ray Excited Emission

Photo-luminescence measured with $\theta = 10^\circ$: No internal absorption



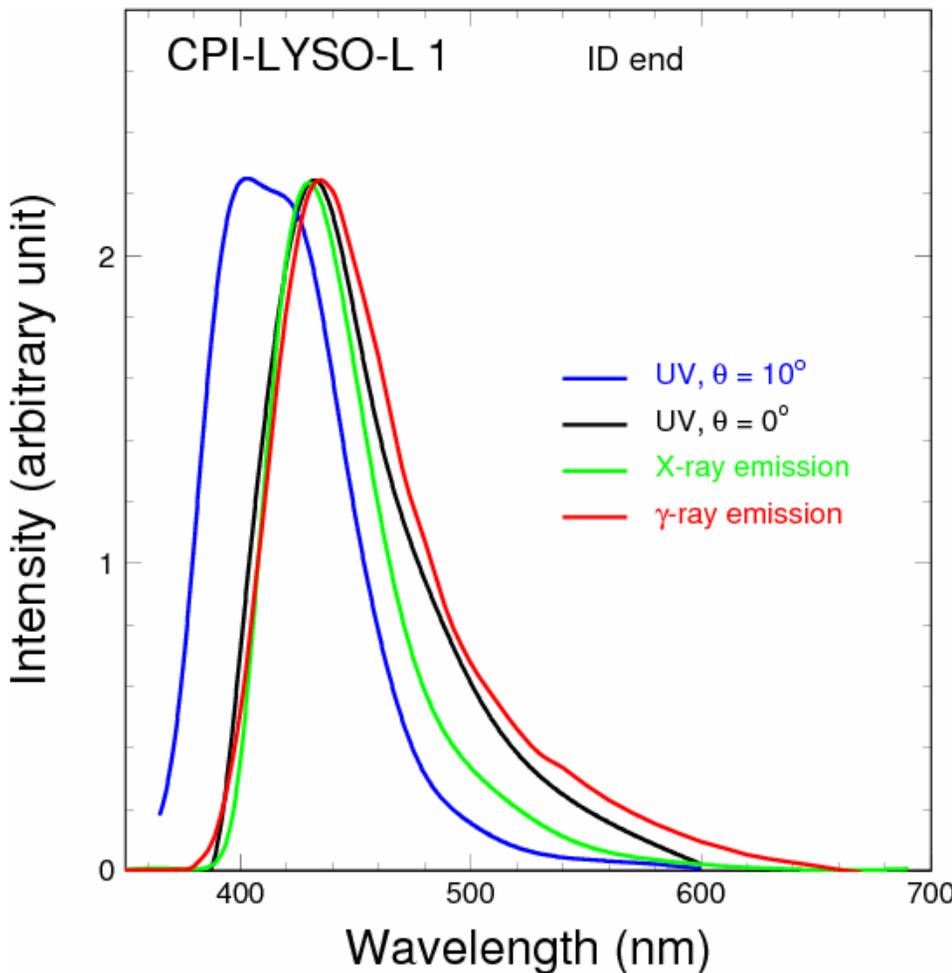
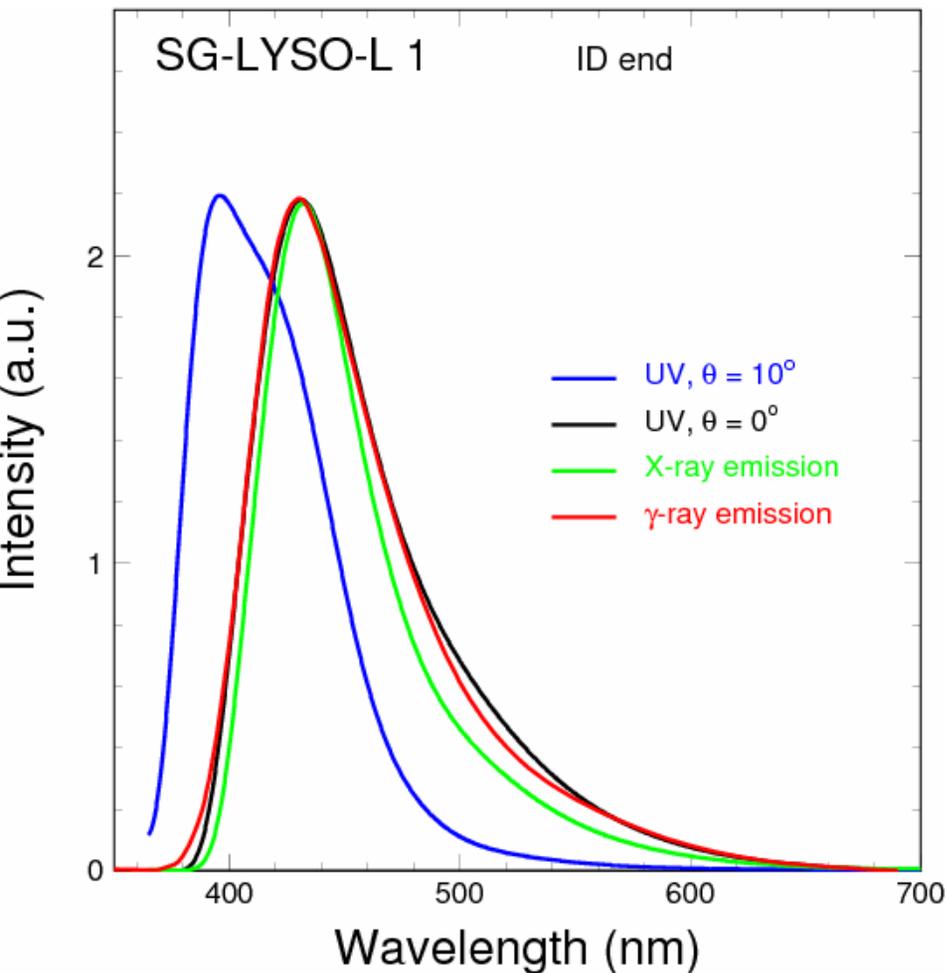
System Checks

UV ($\theta=0^\circ$), X-ray and γ -ray excited emission spectra are consistent
 UV ($\theta=10^\circ$) excited emission has a blue shift because of no absorption



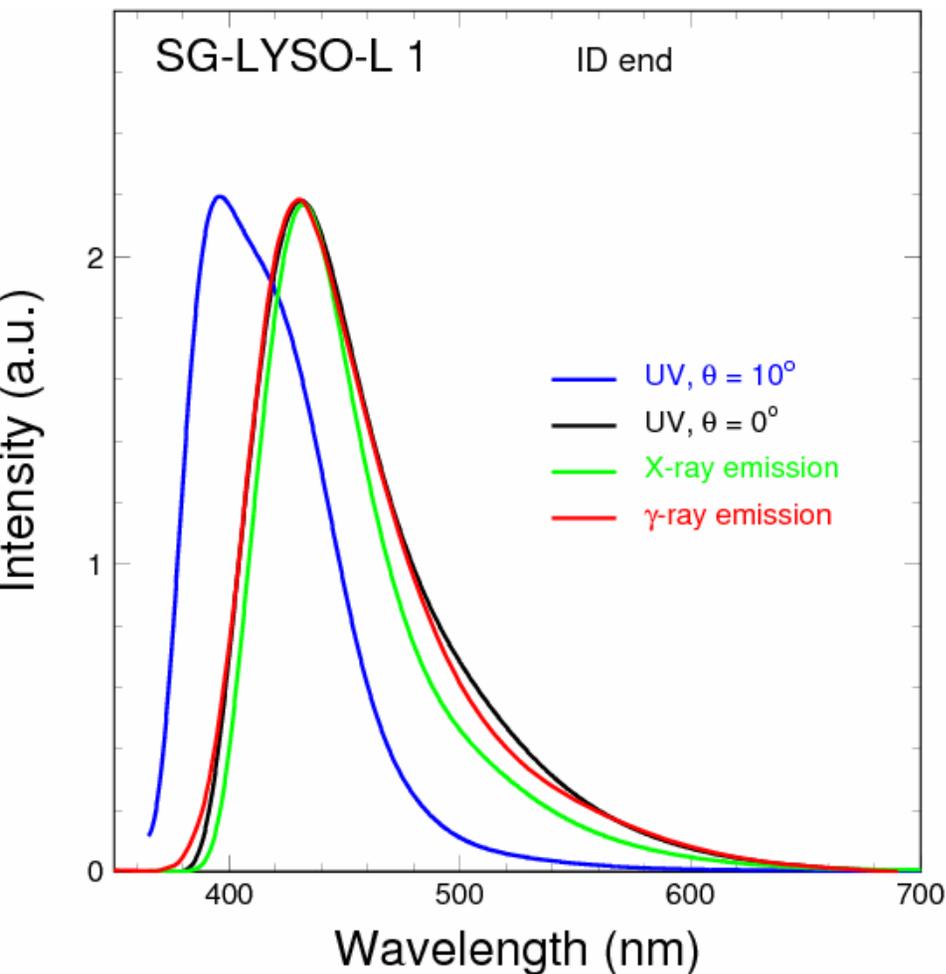
LYSO Emission

UV ($\theta=0^\circ$) and γ -ray excited emission spectra are consistent
 UV ($\theta=10^\circ$) excited emission has a strong blue shift (See N49-1)
 X-ray excited emission is slightly narrow. Why?



X-ray Excited Emission Narrow

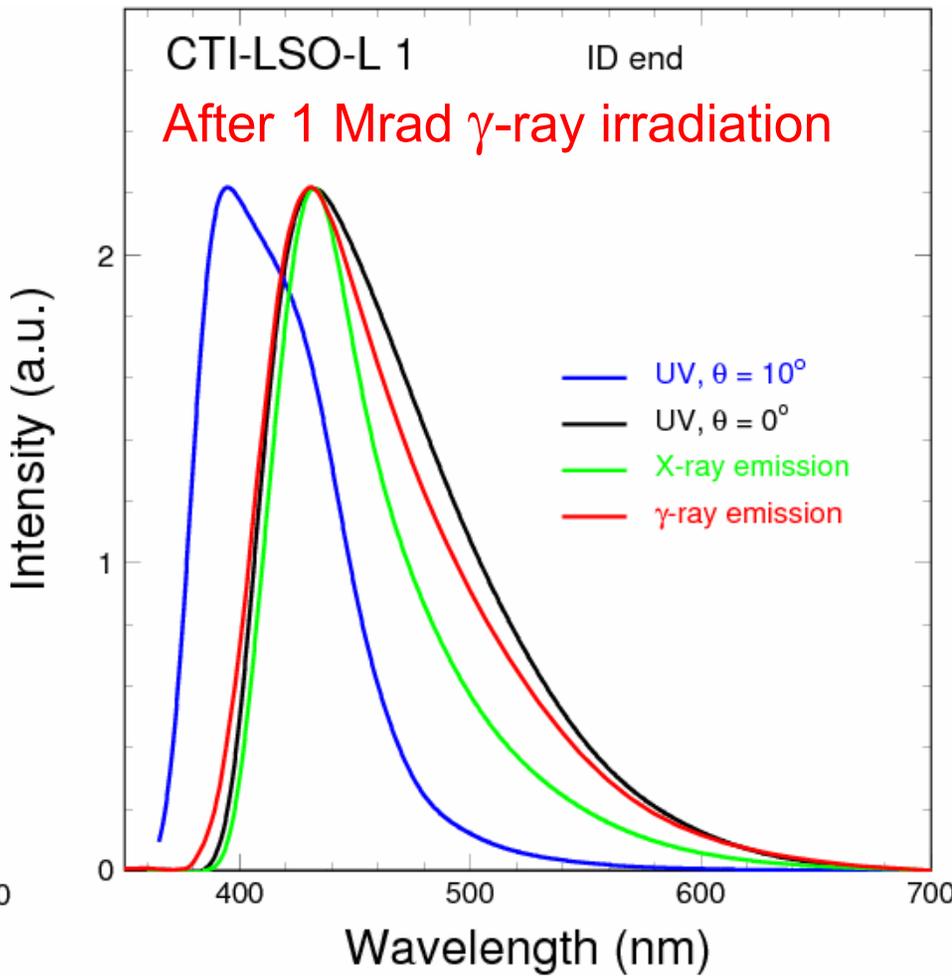
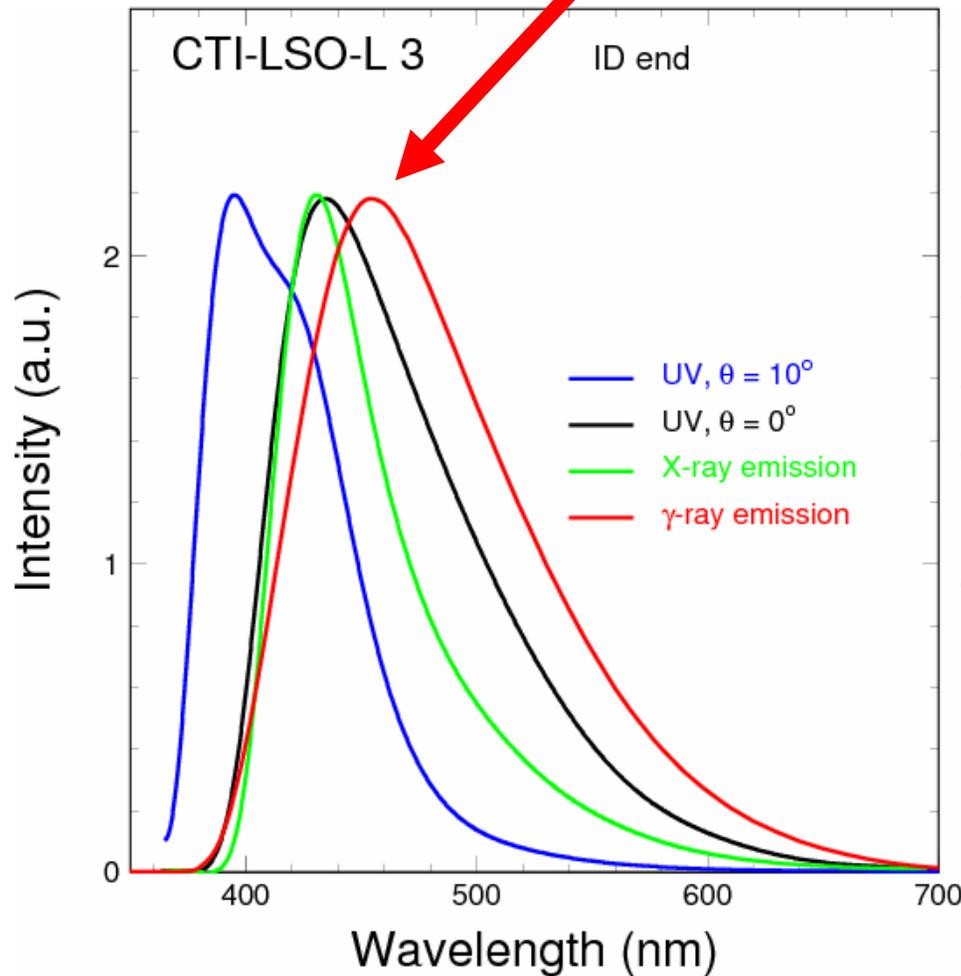
The narrow X-ray excited emission spectra of LYSO may be explained by a surface effect since X-ray does not penetrate.



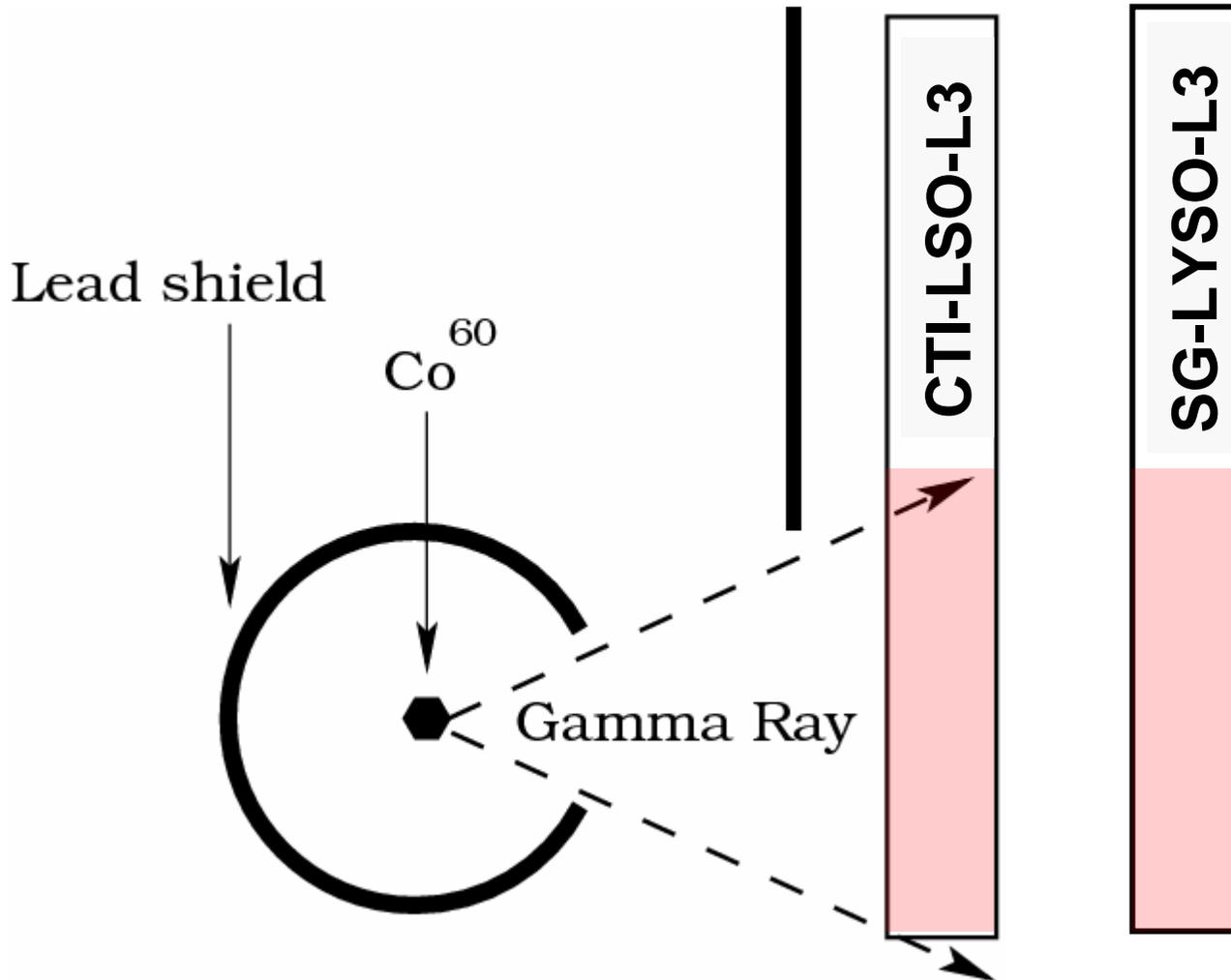
UV excitation (3-6 eV)	Attenuation length 100~700 μm
X-ray (8-30 keV)	Attenuation length $\sim 10 \mu\text{m}$
γ -ray (Co^{60}) ($\sim 1.2 \text{ MeV}$)	Radiation length 1.14 cm

LSO Emission

All emission spectra are similar to that of LYSO, except that γ -ray excited emission has a "red shift", which disappeared after irradiations with γ -ray.



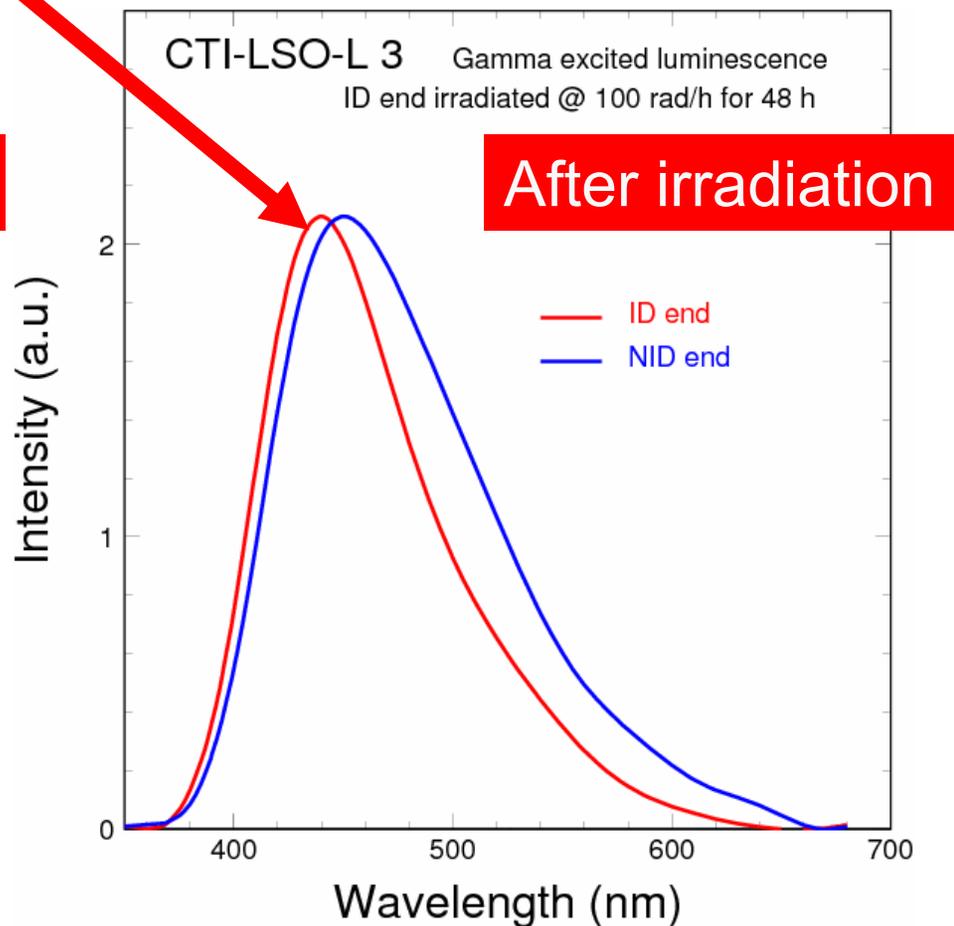
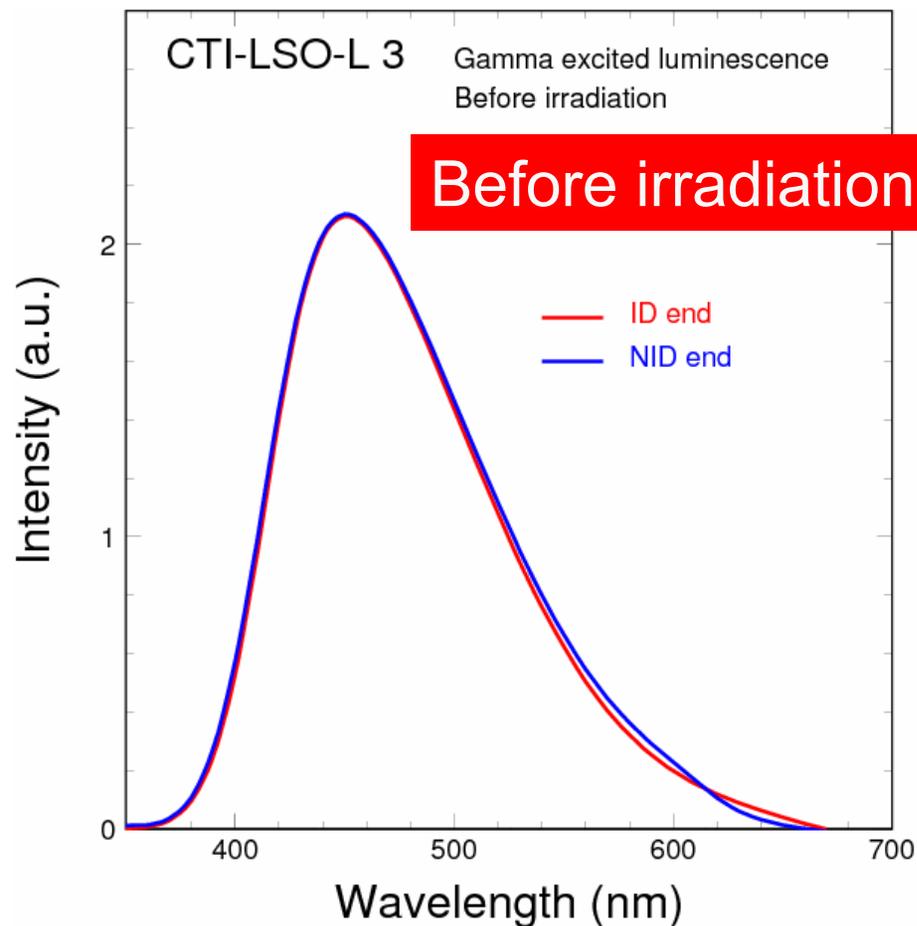
γ -Ray Irradiation on Sample's ID End



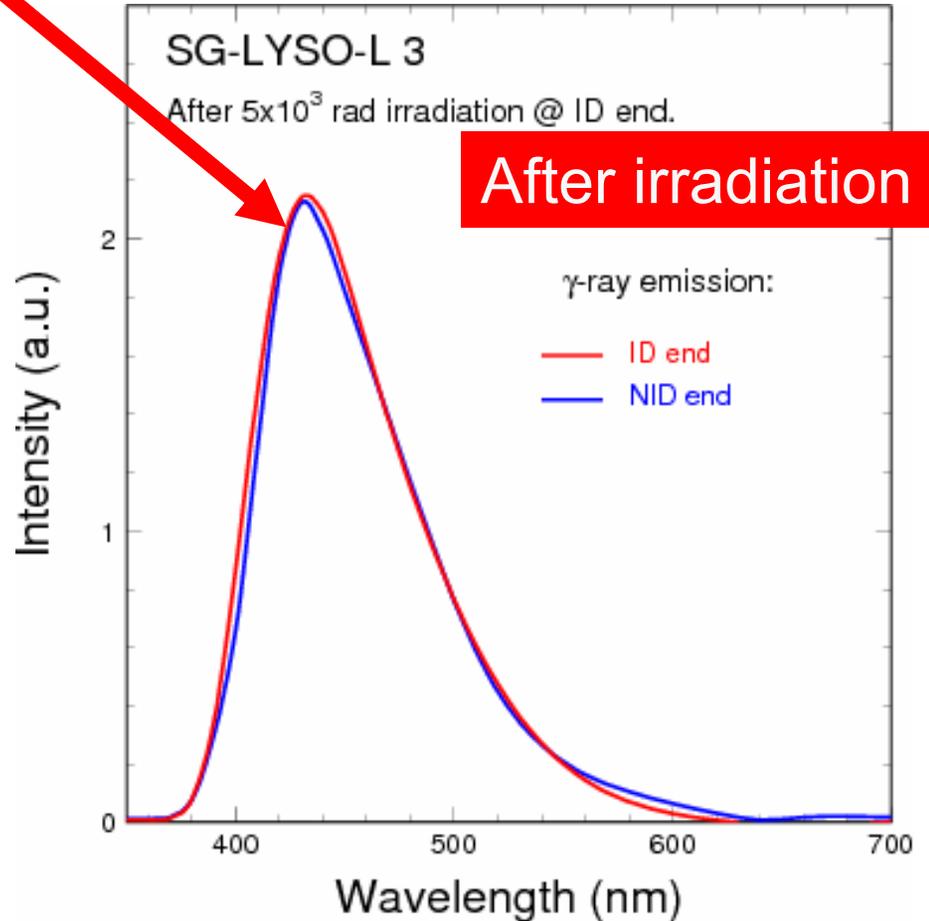
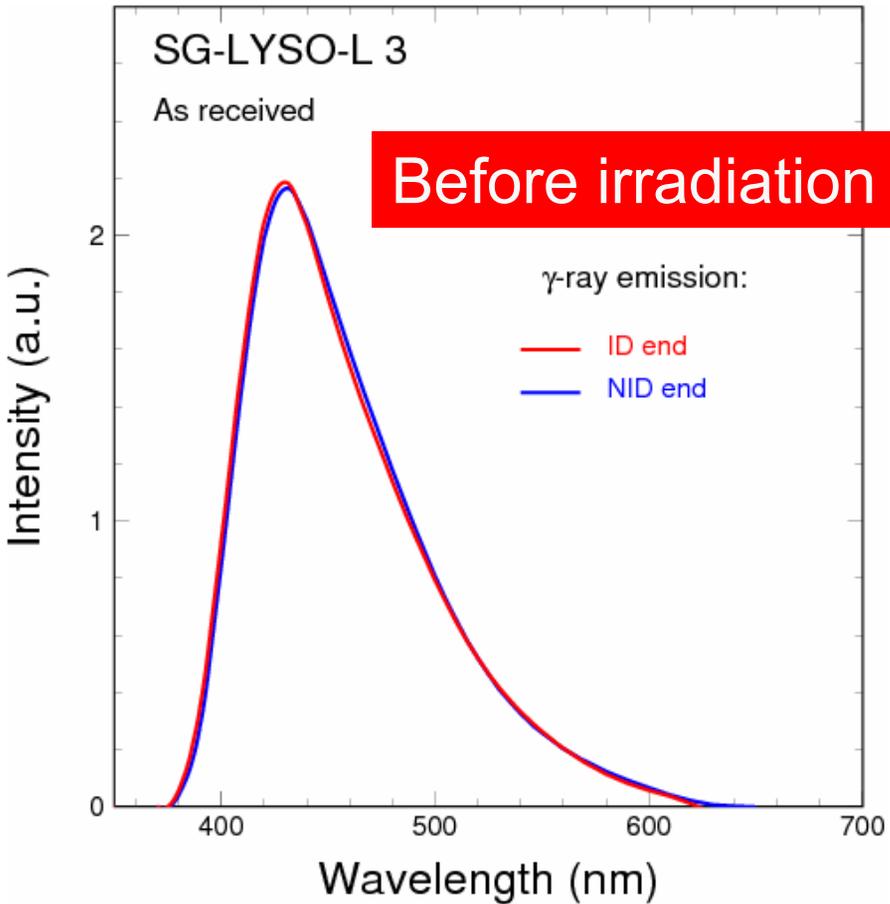
ID End received
~5000 rad

LSO: γ -Ray Excited Emission Spectra

The emission peak of sample's irradiated ID end has a ~ 15 nm "blue" shift

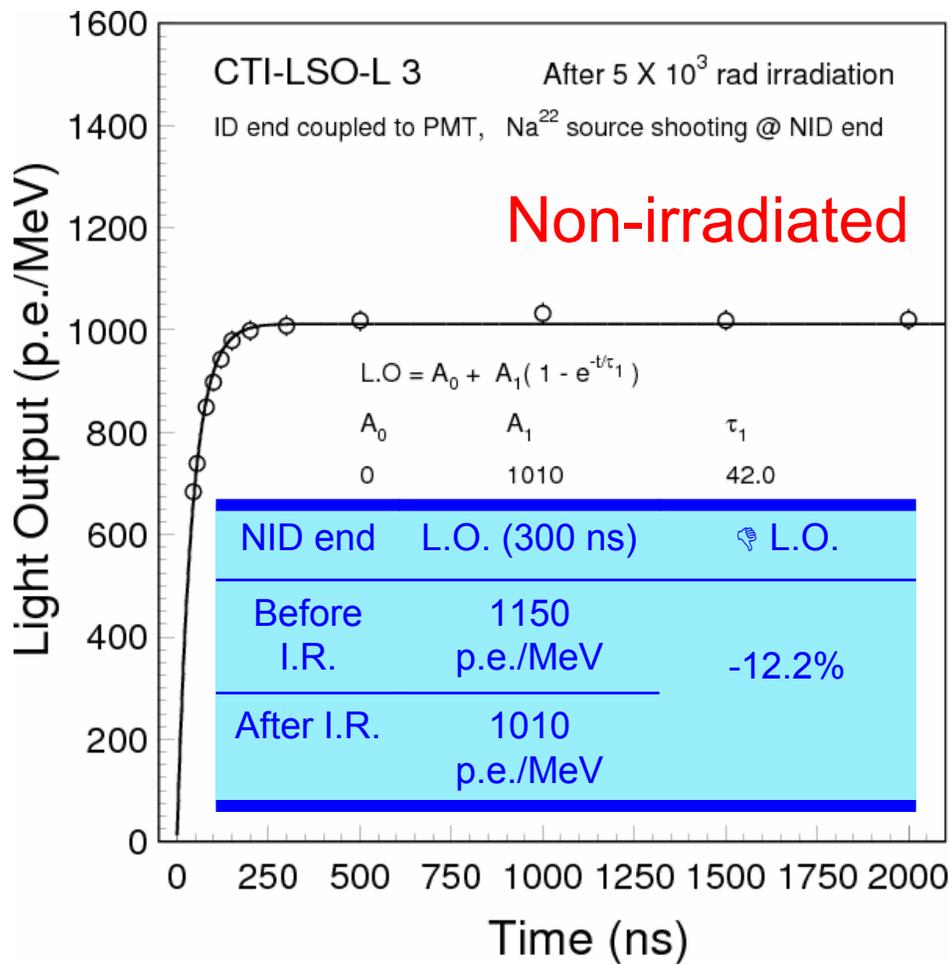
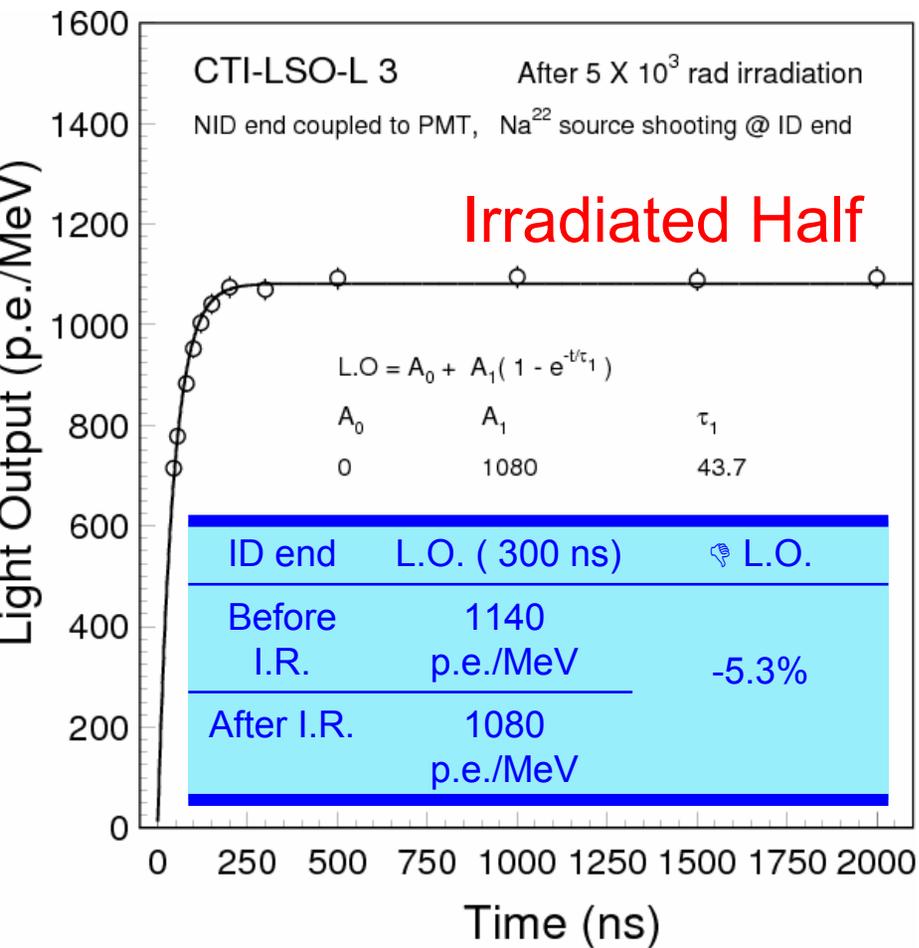


The emission peak of sample's ID (irradiated) end has NO "blue" shift



LSO Decay Time and Light Output

The irradiated end (ID) has no change in decay time. Its light output degradation is half of that of the NID end because of the emission “blue shift”.

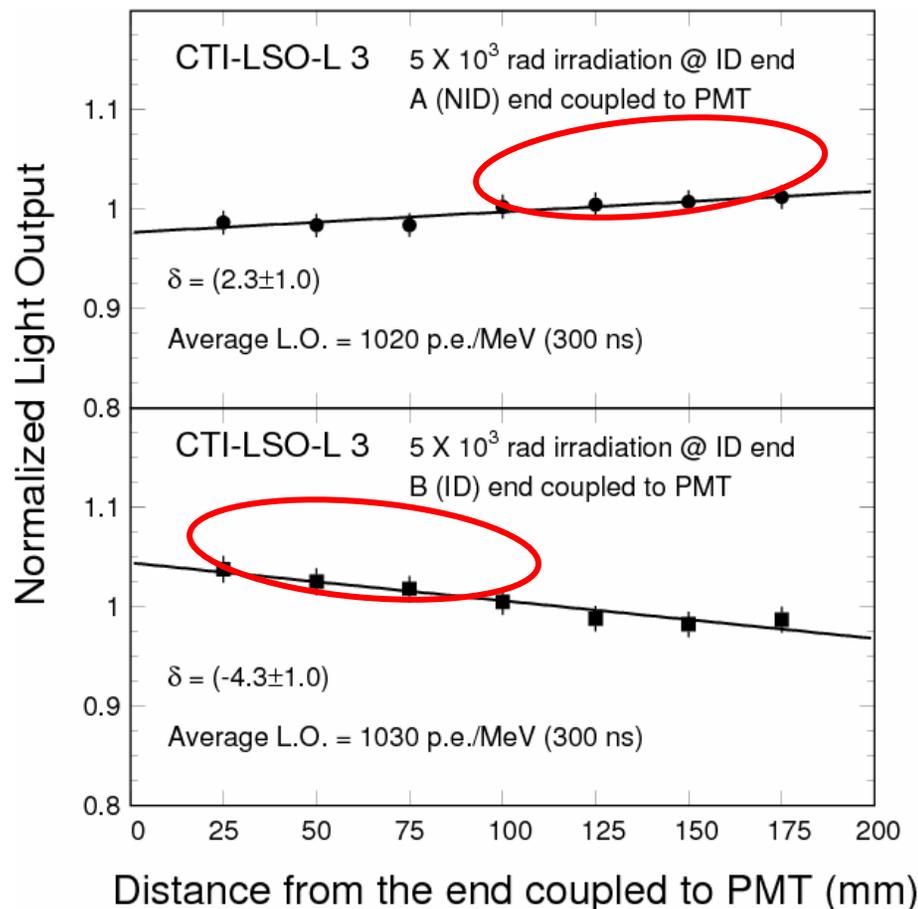
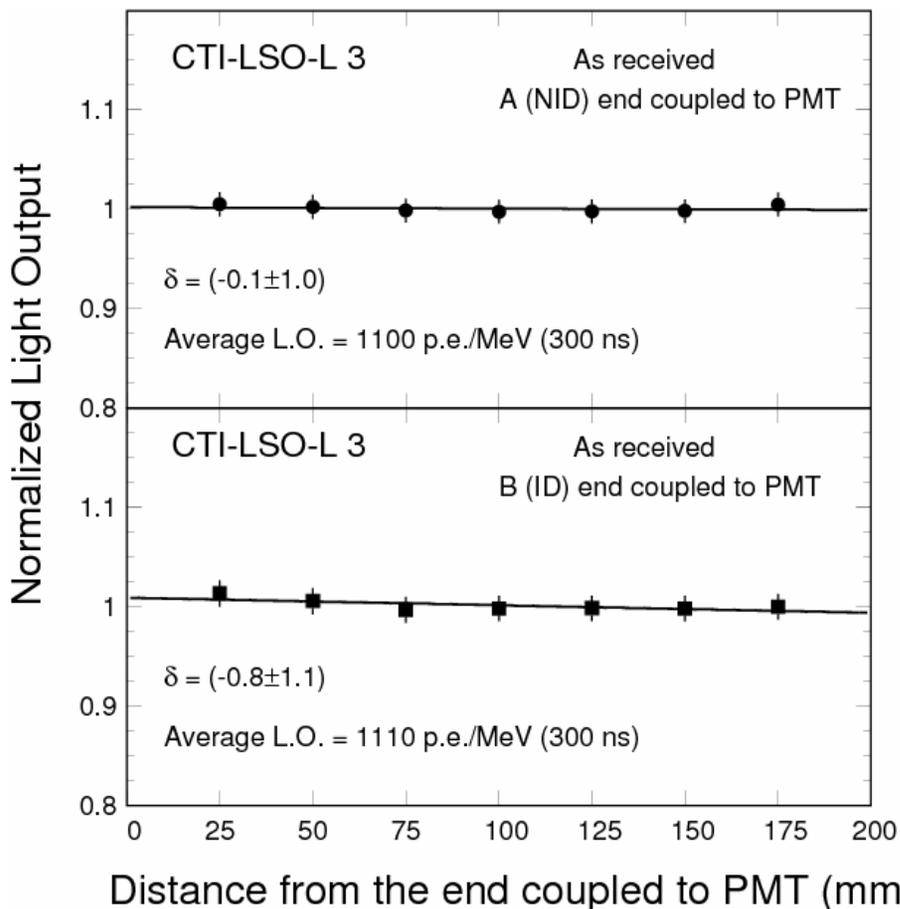


LSO Uniformity with PMT Readout

The emission “blue shift” of the irradiated end causes a relative larger LO for the PMT readout.

Before irradiation

After irradiation

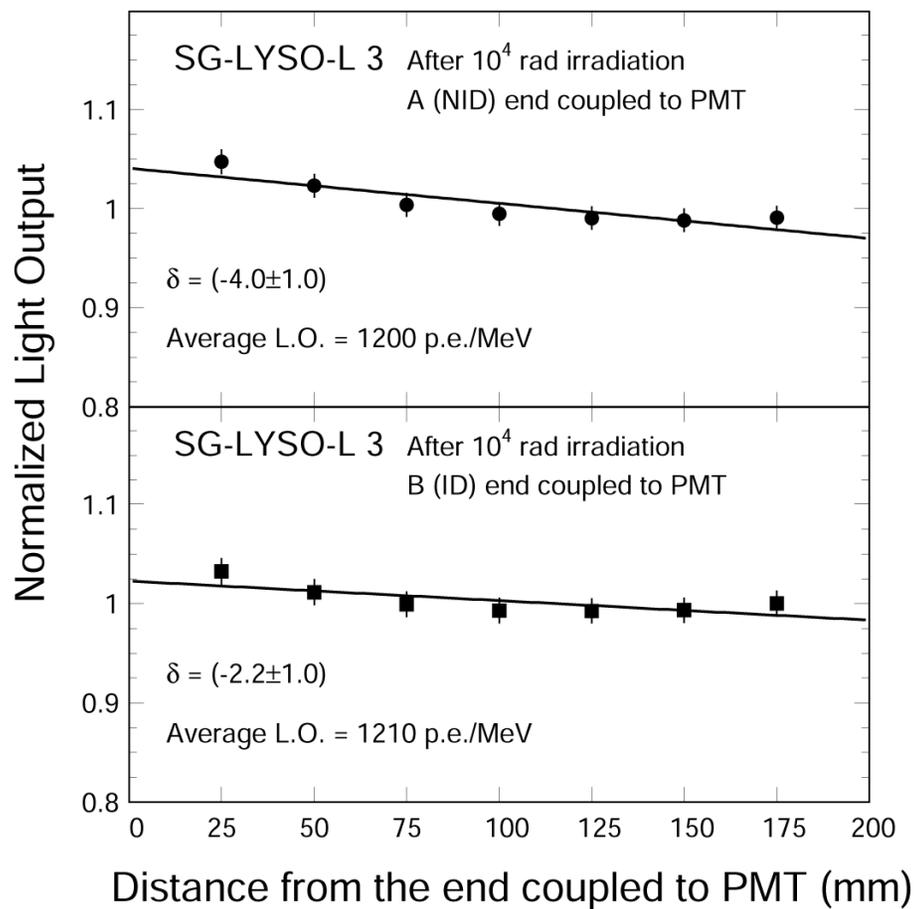
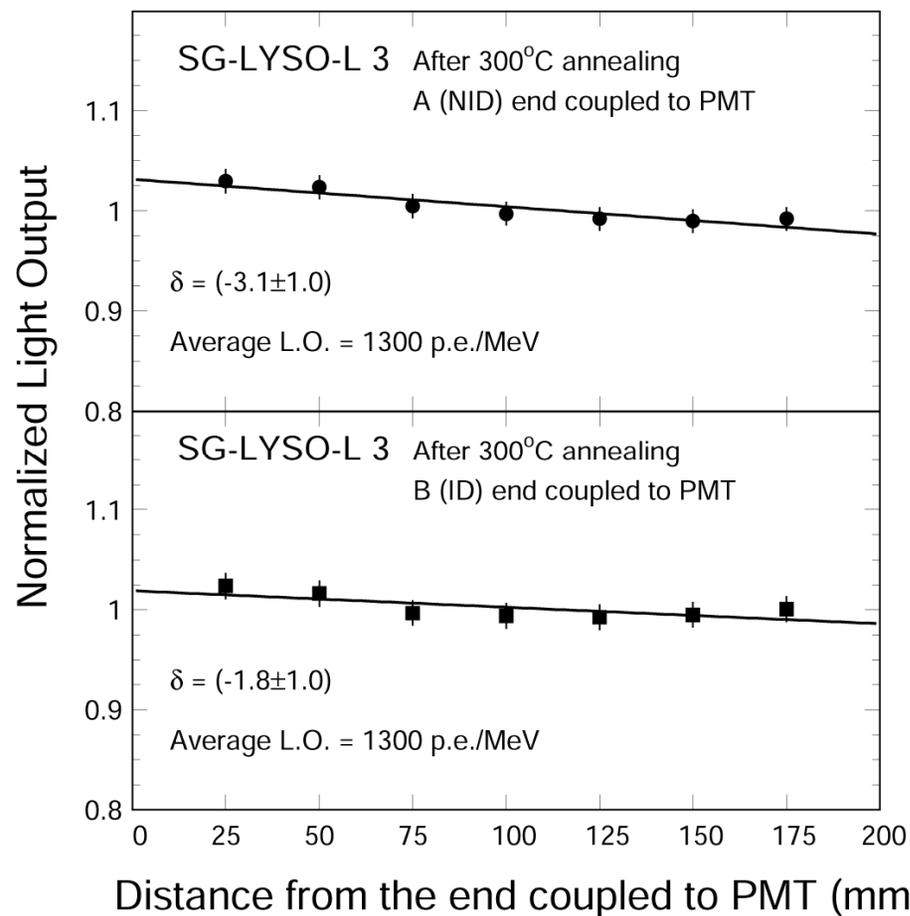


LYSO Uniformity with PMT Readout

No significant variations in the light output and light response uniformity for the PMT readout

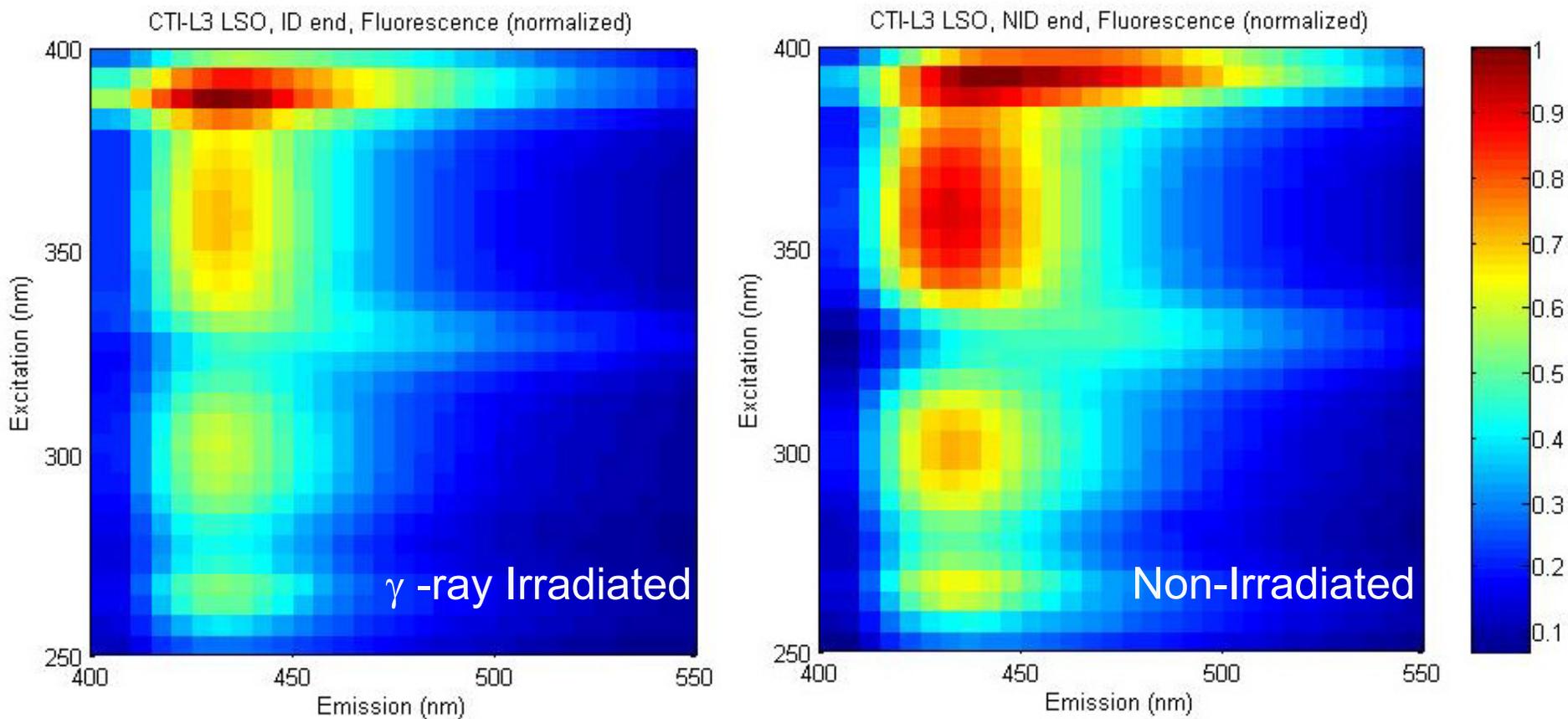
Before irradiation

After irradiation



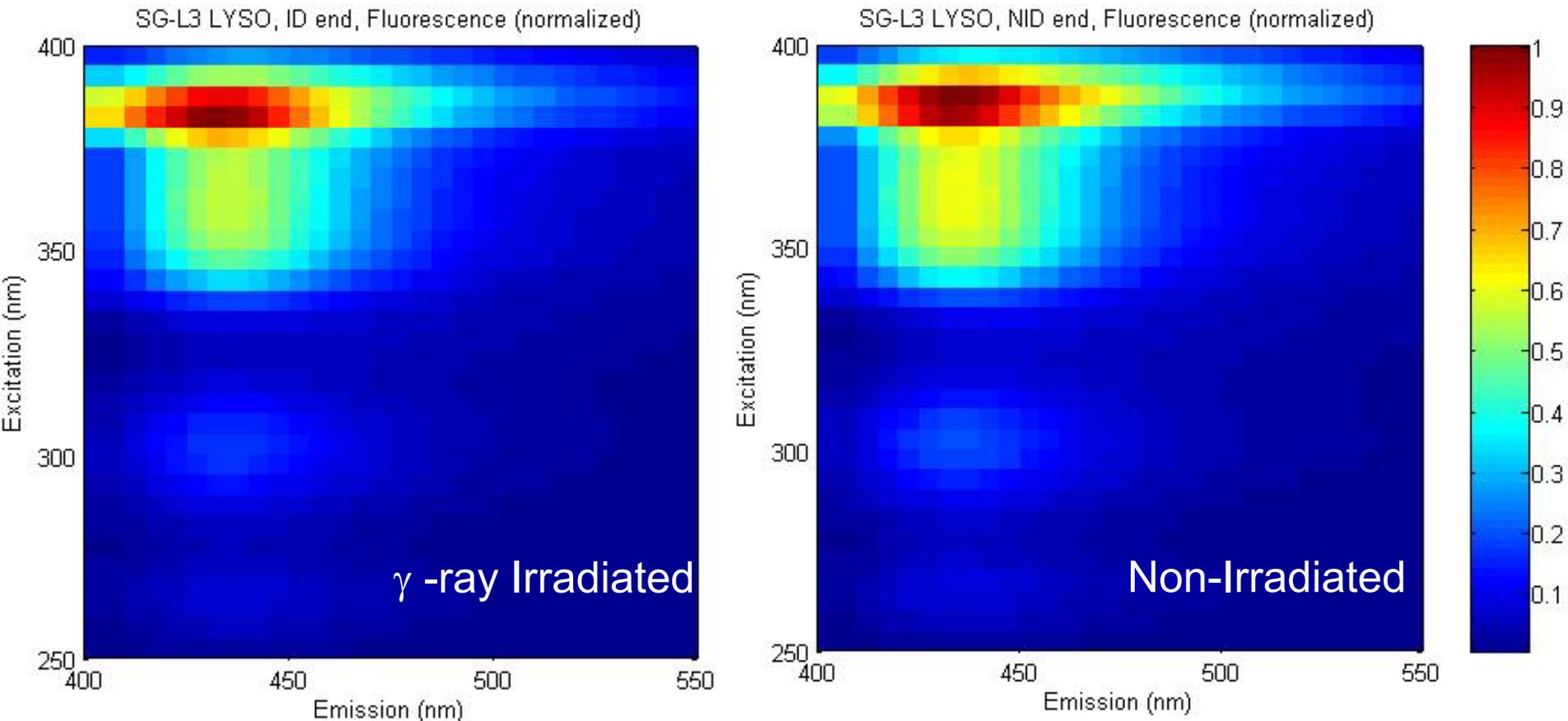
UV Excited Emission Spectra of Two Halves of the LSO Sample

The γ -ray irradiated half shows less long wavelength emission when excited at 325 nm and 380 nm.



UV Excited Emission Spectra of Two Halves of the LYSO Sample

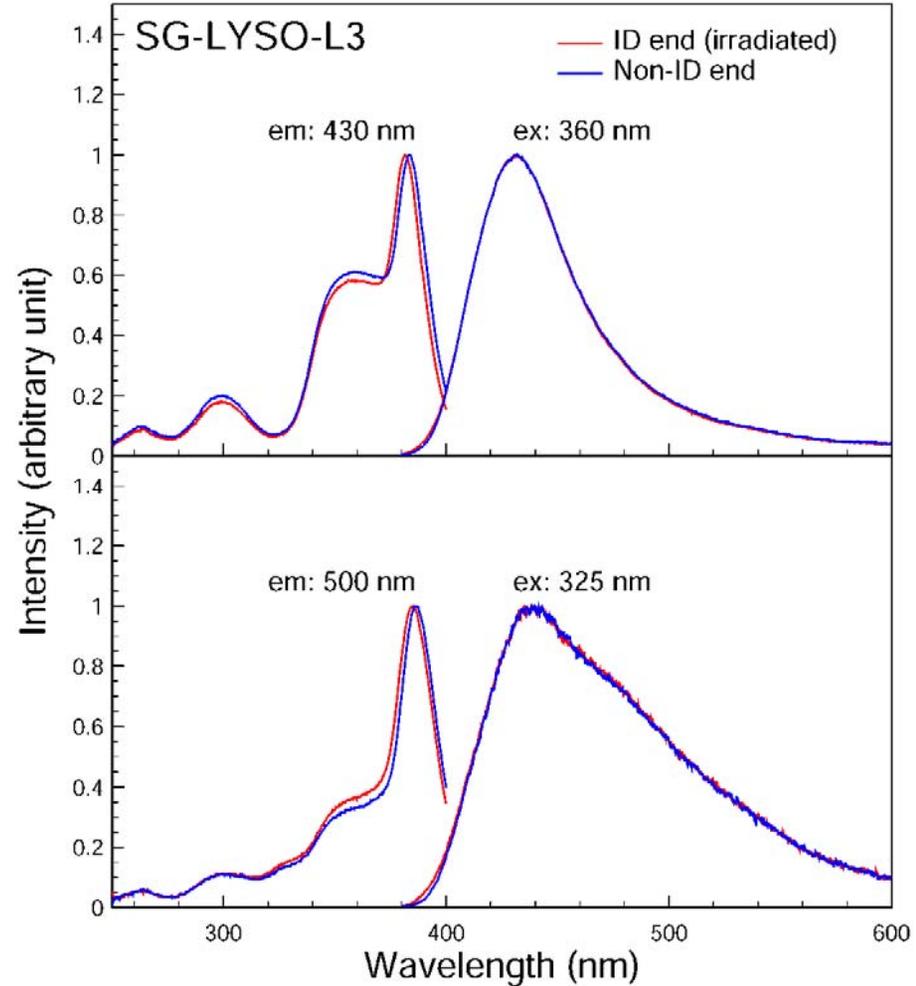
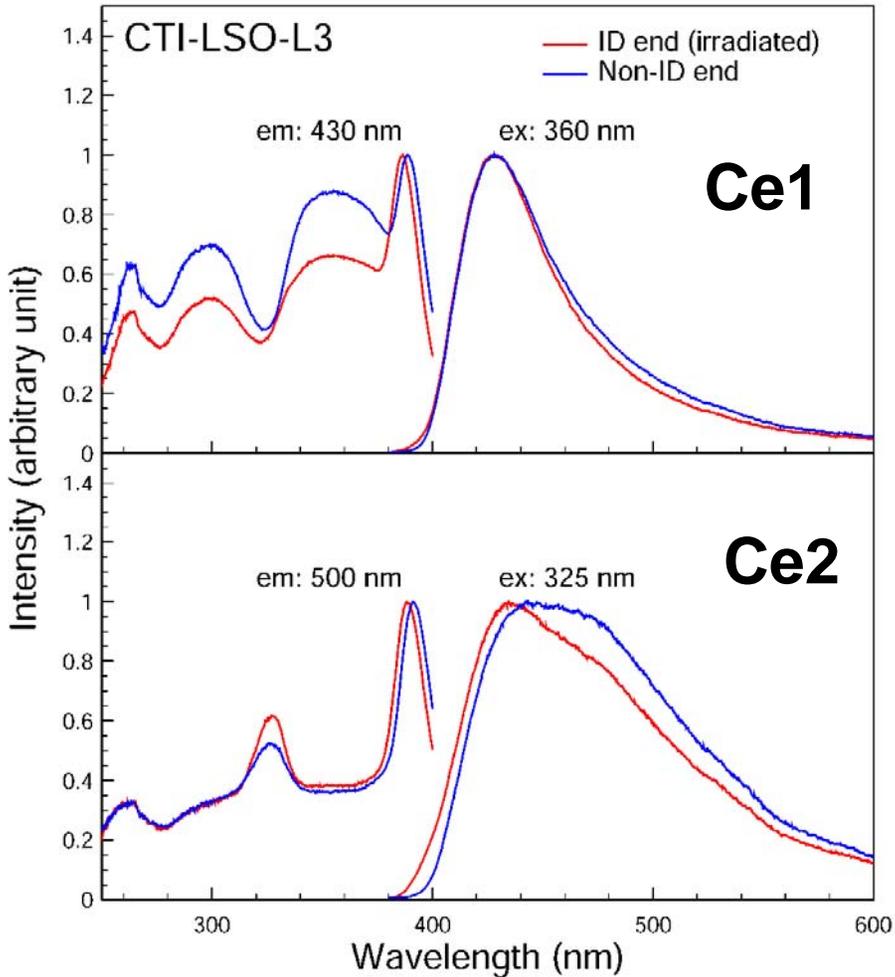
The γ -ray irradiated half shows consistent emission as the non irradiated half when excited at 325 nm and 380 nm.



Ce³⁺ Luminescence Centers in LSO

J.D. Naud et. al., IEEE Trans. Nuclear Sci., Vol.43, p1324, June 1996

Ce1: two regular Lu³⁺ crystallographic sites, ex: 360 nm, em: 430 nm
 Ce2: irregular sites, proposed "interstitial site", ex: 325 nm, em: 500 nm



Conclusions

- A strong blue shift of the photo luminescence ($\theta=10^\circ$) in LSO/LYSO is attributed to its self absorption.
- A narrow X-ray excited emission spectra in LSO/LYSO seems caused by a surface effect
- A broader γ -ray excited emission spectrum with a “red shift” as compared to the X-ray and UV excited emission spectra is observed in large size LSO samples. This shift disappeared after γ -ray irradiations. This observation consists with the light output and uniformity data and with what reported in NSS05 at Puerto Rico.
- No such shift was observed in large size LYSO samples.
- We tentatively attribute this shift to the contribution of the “irregular” sites of Ce^{3+} (the component around 450 nm). The fact that it can be “cured” more or less by γ -ray irradiations supports that this site is a defect perturbed irregular site of Ce^{3+} .